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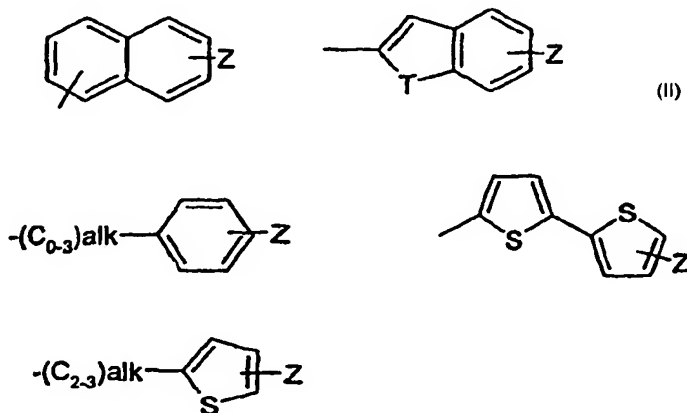
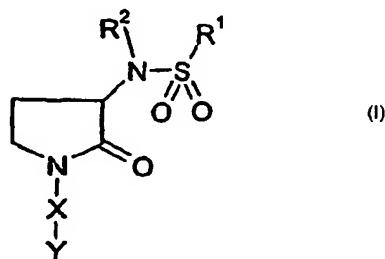
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(54) Title: 1-PHENYL-2-OXO-3-SULFONYLAMINO-PYRROLIDINE DERIVATIVES AND RELATED COMPOUNDS AS
FACTOR XA INHIBITORS FOR THE TREATMENT OF ACUTE VASCULAR DISEASES



(57) Abstract: The invention relates to compounds of formula (I) wherein: R¹ represents a group selected from: formula (II) each ring of which optionally contains a further heteroatom N, Z represents an optional substituent halogen, alk represents alkylene or alkenylene, T represents S, O or NH; R² represents -C₁₋₆alkyl, -C₁₋₃alkylCN, -C₀₋₃alkylR^c, -C₁₋₃alkylR^f, -C₂₋₃alkylNR^aR^b, -C₂₋₃alkylOC₁₋₆alkyl, -C₂₋₃alkylOC₁₋₃alkylCONR^aR^b, with the proviso that R² does not represent C₂₋₃alkylmorpholino; X represents phenyl or a 5- or 6- membered aromatic heterocyclic group containing at least one heteroatom selected from O, N or S, each of which is optionally substituted by 0-2 groups selected from: halogen, -C₁₋₄alkyl, -C₂₋₄alkenyl, -CN, -CF₃, -NR^aR^b, -C₀₋₄alkylOR^c, -C(O)R^d and -C(O)NR^aR^b; Y represents a substituent selected from hydrogen, halogen, -C₁₋₄alkyl, -C₂₋₄alkenyl, -NR^aR^b, -NO₂, -C(O)NR^aR^b, -N(C₁₋₄alkyl)(CHO), -NHCOC₁₋₄alkyl, -NHCO₂R^d, -C₀₋₄alkylOR^c, -C(O)R^d, -S(O)_nR^d, or -S(O)₂NR^aR^b; The other substituents are as defined in claim 1.



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